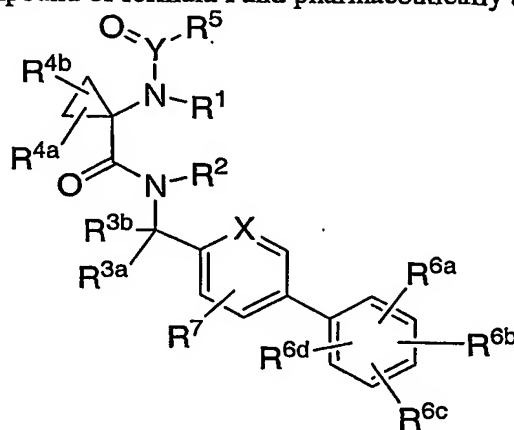


## WHAT IS CLAIMED IS:

1. A compound of formula I and pharmaceutically acceptable salts thereof:



I

wherein

R<sup>1</sup> and R<sup>2</sup> are independently selected from hydrogen and C<sub>1-4</sub> alkyl;

R<sup>3a</sup> and R<sup>3b</sup> are independently selected from hydrogen and C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms;

- 10 R<sup>4a</sup> and R<sup>4b</sup> are independently selected from hydrogen, halogen, and C<sub>1-4</sub> alkyl optionally substituted with 1 to 4 groups selected from halogen, OR<sup>a</sup>, OC(O)R<sup>a</sup>, S(O)<sub>k</sub>R<sup>d</sup>, OS(O)<sub>2</sub>R<sup>d</sup>, and NR<sup>1</sup>R<sup>2</sup>, or R<sup>4a</sup> and R<sup>4b</sup> together with the carbon atom to which they are both attached form an exo-cyclic methylene optionally substituted with 1 to 2 groups selected from C<sub>1-4</sub> alkyl optionally substituted with 1-5 halogens and C<sub>1-4</sub> alkyloxy;

- 15 R<sup>5</sup> is selected from (1) C<sub>1-6</sub> alkyl optionally substituted with 1 to 5 groups independently selected from halogen, nitro, cyano, OR<sup>a</sup>, SR<sup>a</sup>, COR<sup>a</sup>, SO<sub>2</sub>R<sup>d</sup>, CO<sub>2</sub>R<sup>a</sup>, OC(O)R<sup>a</sup>, NR<sup>b</sup>R<sup>c</sup>, NR<sup>b</sup>C(O)R<sup>a</sup>, NR<sup>b</sup>C(O)<sub>2</sub>R<sup>a</sup>, C(O)NR<sup>b</sup>R<sup>c</sup>, C<sub>3-8</sub> cycloalkyl, (2) C<sub>3-8</sub> cycloalkyl optionally substituted with 1 to 5 groups independently selected from halogen, nitro, cyano and phenyl, (3) C<sub>3-6</sub> alkynyl, (4) C<sub>2-6</sub> alkenyl optionally substituted with hydroxyethyl, (5) (CH<sub>2</sub>)<sub>k</sub>-aryl optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, OR<sup>a</sup>, SR<sup>a</sup>, C(O)<sub>2</sub>R<sup>a</sup>, C<sub>1-4</sub> alkyl and C<sub>1-3</sub> haloalkyl; 20 (6) (CH<sub>2</sub>)<sub>k</sub>-heterocycle optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, OR<sup>a</sup>, SR<sup>a</sup>, C<sub>1-4</sub> alkyl and C<sub>1-3</sub> haloalkyl wherein said heterocycle is selected from (a) a 5-membered heteroaromatic ring having a ring heteroatom selected from N, O and S, and optionally having up to 3 additional ring nitrogen atoms wherein said ring is optionally benzo-fused; (b) a 6-membered heteroaromatic ring containing from 1 to 3 ring nitrogen atoms and N-oxides thereof, wherein said ring is 25 optionally benzo-fused; and (c) a 5- or 6-membered non-aromatic heterocyclic ring selected from

tetrahydrofuranyl, 5-oxotetrahydrofuranyl, 2-oxo-2H-pyranyl, 6-oxo-1,6-dihydropyridazinyl, (7) C(O)<sub>2</sub>R<sup>a</sup>, and (8) C(O)NR<sup>b</sup>R<sup>c</sup>;

R<sup>6a</sup> is selected from (1) -OSO<sub>2</sub>R<sup>8</sup>, (2) -NR<sup>8a</sup>SO<sub>2</sub>R<sup>9</sup>, and (3) -C(R<sup>8b</sup>)(R<sup>8c</sup>)SO<sub>2</sub>R<sup>9</sup>;

R<sup>6b</sup>, R<sup>6c</sup>, and R<sup>6d</sup> are independently selected from (1) hydrogen, (2) halogen, (3) OSO<sub>2</sub>R<sup>8</sup>, (4) C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms, (5) cyano, (6) nitro, (7) OR<sup>a</sup>, and (8) CO<sub>2</sub>R<sup>a</sup>, or when attached to adjacent carbon atoms R<sup>6c</sup> and R<sup>6d</sup> together with the carbon atoms to which they are attached form a 5- to 8-membered saturated or unsaturated ring;

R<sup>7</sup> is selected from (1) hydrogen, (2) halogen, (3) cyano, (4) nitro, (5) OR<sup>a</sup>, (6) CO<sub>2</sub>R<sup>a</sup>, (7) C(O)NR<sup>b</sup>R<sup>c</sup>, and (8) C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms,

R<sup>8</sup> is selected from (1) C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms, (2) (CH<sub>2</sub>)<sub>k</sub>-aryl optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, NR<sup>a</sup>C(O)R<sup>a</sup>, OR<sup>a</sup>, SR<sup>a</sup>, CO<sub>2</sub>R<sup>a</sup>, C<sub>1-4</sub> alkyl, C<sub>1-3</sub> haloalkyl and NR<sup>b</sup>R<sup>c</sup>, (3) NR<sup>b</sup>R<sup>c</sup>, and (4) hydrogen;

R<sup>8a</sup> is selected from hydrogen, C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms, halogen, and CO<sub>2</sub>R<sup>a</sup>, or

when R<sup>6a</sup> and R<sup>6b</sup> are attached to adjacent atoms, R<sup>8a</sup> and R<sup>6b</sup> together complete 5- or 6-membered ring;

R<sup>8b</sup> and R<sup>8c</sup> are independently selected from hydrogen, C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms, halogen, cyano, nitro, CO<sub>2</sub>R<sup>a</sup>, and OR<sup>a</sup>;

R<sup>9</sup> is selected from (1) C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms, (2) aryl optionally

substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, NR<sup>a</sup>C(O)R<sup>a</sup>, OR<sup>a</sup>, SR<sup>a</sup>, CO<sub>2</sub>R<sup>a</sup>, C<sub>1-4</sub> alkyl and C<sub>1-3</sub> haloalkyl, and (3) (CH<sub>2</sub>)<sub>k</sub>-aryl optionally substituted with 1 to 3 groups independently selected from halogen, nitro, cyano, NR<sup>a</sup>C(O)R<sup>a</sup>, OR<sup>a</sup>, SR<sup>a</sup>, C(O)<sub>2</sub>R<sup>a</sup>, C<sub>1-4</sub> alkyl and C<sub>1-3</sub> haloalkyl, or

R<sup>8a</sup> and R<sup>9</sup> together with the atoms to which they are attached form a 5- to 8-membered heterocyclic ring;

R<sup>a</sup>, R<sup>b</sup> and R<sup>c</sup> are independently selected from (1) hydrogen, (2) C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms, (3) phenyl optionally substituted with 1 to 3 groups selected from halogen, cyano, nitro, OH, C<sub>1-4</sub> alkyloxy, C<sub>3-6</sub> cycloalkyl and C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms, and (4) C<sub>3-6</sub> cycloalkyl, or

R<sup>b</sup> and R<sup>c</sup> together with the nitrogen atom to which they are attached form a 4-, 5-, or 6-membered ring optionally containing an additional heteroatom selected from N, O, and S; or

R<sup>b</sup> and R<sup>c</sup> together with the nitrogen atom to which they are attached form a cyclic imide;

R<sup>d</sup> is selected from (1) C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms, (2) C<sub>1-4</sub> alkyloxy, (3) phenyl optionally substituted with 1 to 3 groups selected from halogen, cyano, nitro, OH, C<sub>1-4</sub>

alkyloxy, C<sub>3-6</sub> cycloalkyl and C<sub>1-4</sub> alkyl optionally substituted with 1 to 5 halogen atoms, and (4) hydrogen;

X is selected from CH and N;

Y is selected from C and S=O; and

5 k is selected from 0, 1, and 2.

2. A compound of Claim 1 wherein R<sup>5</sup> is selected from pyrimidinyl and C<sub>1-6</sub> alkyl optionally substituted with 1 to 5 groups independently selected from halogen.

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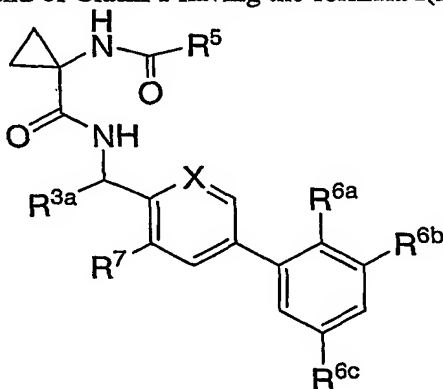
3. A compound of Claim 1 wherein Y is C.

4. A compound of Claim 1 wherein R<sup>6a</sup> is OSO<sub>2</sub>R<sup>8</sup> and R<sup>8</sup> is selected from 2,2,2-trifluoroethyl, trifluoromethyl, methyl, ethyl, propyl, isopropyl, phenyl, benzyl, and dimethylamino; or  
15 R<sup>6a</sup> is NHSO<sub>2</sub>R<sup>9</sup> and R<sup>9</sup> is methyl or trifluoromethyl.

5. A compound of Claim 1 wherein R<sup>6b</sup> is selected from hydrogen, fluorine, and chlorine.

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6. A compound of Claim 1 having the formula I(2):



I(2)

wherein X is N or CH, R<sup>3a</sup> is H or C<sub>1-4</sub>alkyl, R<sup>7</sup> is hydrogen or halogen, and R<sup>5</sup>, R<sup>6a</sup>, R<sup>6b</sup> and R<sup>6c</sup> have the same definitions as provided in Claim 1.

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7. A compound of Claim 6 wherein R<sup>6a</sup> is NHSO<sub>2</sub>R<sup>9</sup>; R<sup>9</sup> is C<sub>1-4</sub>alkyl optionally substituted with 1 to 5 halogen atoms, R<sup>6b</sup> is halogen, and R<sup>6c</sup> is hydrogen or halogen.

8. A compound of Claim 10 wherein R<sup>6a</sup> is OSO<sub>2</sub>R<sup>8</sup>; R<sup>8</sup> is selected from methyl, trifluoromethyl, ethyl, propyl, isopropyl, benzyl, dimethylamino, 2,2,2-trifluoroethyl, and phenyl; R<sup>6b</sup> is hydrogen or halogen, and R<sup>6c</sup> is hydrogen or halogen.

9. A compound of Claim 10 wherein R<sup>5</sup> is pyrimidinyl or C<sub>1-4</sub>alkyl optionally substituted with 1 to 5 groups independently selected from halogen.

10. A compound selected from  
 3,3'-difluoro-4'-{[(1-[(pyrimidin-5-ylcarbonyl)amino]cyclopropyl)carbonyl]amino}methyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,  
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,  
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoromethyl)sulfonyl]amino}cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,  
 1-[(1R)-1-(3,3'-difluoro-2'-[(trifluoromethyl)sulfonyl]oxy)-1,1'-biphenyl-4-yl]ethyl]amino}carbonyl)-cyclopropanaminium trifluoroacetate,  
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl methanesulfonate,  
 5-chloro-3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,  
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl ethanesulfonate,  
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl propane-1-sulfonate  
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl propane-2-sulfonate,  
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl benzenesulfonate,  
 3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl]amino}ethyl}-1,1'-biphenyl-2-yl phenylmethanesulfonate

3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl)amino]ethyl}-1,1'-biphenyl-2-yl dimethylsulfamate,

3,3'-difluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl)amino]ethyl}-1,1'-biphenyl-2-yl 2,2,2-trifluoroethanesulfonate,

5 3-chloro-3'-fluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl)amino]ethyl}-1,1'-biphenyl-2-yl trifluoromethanesulfonate,

3'-fluoro-4'-{(1R)-1-[(1-[(trifluoroacetyl)amino]cyclopropyl)carbonyl)amino]ethyl}-2-[(trifluoromethyl)sulfonyl]oxy}-1,1'-biphenyl-3-yl trifluoromethanesulfonate,

10 N-(1-[(1-[(1R)-1-(3,3'-difluoro-2'-[methyl(methylsulfonyl)amino]-1,1'-biphenyl-4-yl)ethyl)amino]carbonyl]cyclopropyl)pyrimidine-5-carboxamide,

N-(1-[(1-[(3,3'-difluoro-2'-[(methylsulfonyl)amino]-1,1'-biphenyl-4-yl)methyl)amino]carbonyl]cyclopropyl)pyrimidine-5-carboxamide,

N-{1-[(1-[(2'-(1,1-dioxido-1,2-thiazinan-2-yl)-3,3'-difluoro-1,1'-biphenyl-4-yl)methyl]amino)carbonyl]cyclopropyl}pyrimidine-5-carboxamide,

15 N-[(1R)-1-(3,3'-difluoro-2'-[(trifluoromethyl)sulfonyl]methyl)-1,1'-biphenyl-4-yl)ethyl]-1-[(trifluoroacetyl)amino]cyclopropanecarboxamide,

N-[(1R)-1-(3,3'-difluoro-2'-[(trifluoromethyl)sulfonyl]amino)-1,1'-biphenyl-4-yl)ethyl]-1-[(trifluoroacetyl)amino]cyclopropanecarboxamide, and

20 N-(1-[(1-[(1R)-1-(3,3'-difluoro-2'-[(methylsulfonyl)amino]-1,1'-biphenyl-4-yl)ethyl)amino]carbonyl]cyclopropyl)pyrimidine-5-carboxamide.

11. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1 and a pharmaceutically acceptable carrier.

25 12. Use of a compound of Claim 1 or a pharmaceutically acceptable salt thereof in the manufacture of a medicament for the treatment or prevention of pain and inflammation.

13. Use of Claim 30 wherein said pain is postherpetic neuropathy, osteoarthritis pain, or dental pain.

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